**Request to Emory IRB for Determination of Sensitive Status**

**IRB File #**

**PI:**

**Date:**

Instructions: Investigators should complete this form and upload it in the eIRB smartform (Data and Safety Monitoring section, 6.0) to request “sensitive study” status for a clinical study.

**Sensitive studies are subject to the following processes:**

* The Office of Clinical Research (OCR) will NOT link the signed consent/HIPAA documents in the EeMR
* OCR will link the *Sensitive Study version of the Key Points Summary* to the EeMr. This version has key safety information but not the study title.

Please answer all of the following questions**:**

1. **What is the stigmatizing condition or factor?** (Note: Illicit drug use, behaviors placing a subject at high risk of HIV or hepatitis infection, HIV serostatus, hepatitis infection, schizophrenia, and bipolar disorder are presumed to be stigmatizing.)
2. **How is this condition or factor noted in the study (e.g. study title)?**
3. **Will identifiable information about the subject be recorded that could be linked to the specific condition or factor? If so, what information and how will it be linked?**
4. **What other measures are in place to protect the privacy of the subject and the confidentiality of their information?**